

Biomimetic hybrid nanoconstructs for cancer therapy

Original

Biomimetic hybrid nanoconstructs for cancer therapy / Susa, F.; Dumontel, B.; Canta, M.; Racca, L.; Garino, N.; Limongi, T.; Chiodoni, A.; Cauda, V.. - STAMPA. - (2018), pp. 191-191. (Intervento presentato al convegno Merck and Elsevier Young Chemists Symposium (MEYCS 2018) tenutosi a Rimini (italy) nel November 19th – 21st, 2018).

Availability:

This version is available at: 11583/2719168 since: 2018-11-30T16:04:26Z

Publisher:

F. Bella, L. Botta, R. Cucciniello, A. D'Urso, P. Franco, E. Lenci, G. Mazzone, M. Schlich, A. Soldà, R.

Published

DOI:

Terms of use:

openAccess

This article is made available under terms and conditions as specified in the corresponding bibliographic description in the repository

Publisher copyright

(Article begins on next page)



MERCK & ELSEVIER Young Chemists Symposium



An event by



Società Chimica Italiana
Gruppo Giovani



BOOK OF ABSTRACTS

RIMINI (ITALY)
NOVEMBER 19TH – 21ST, 2018



ELSEVIER

MERCK

Proceedings of the
Merck & Elsevier Young Chemists Symposium
XVIII edition

Edited by: F. Bella, L. Botta, R. Cucciniello, A. D'Urso, P. Franco, E. Lenci, G. Mazzone, M. Schlich, A. Soldà, R. Spezzano, S. Staderini, and L. Triggiani

Copyright © 2018 Società Chimica Italiana, Viale Liegi 48C, 00198-Roma

ISBN: 978-88-94952-03-2

Biomimetic hybrid nanoconstructs for cancer therapy

Francesca Susa,^a Bianca Dumontel,^a Marta Canta,^a Luisa Racca,^a Nadia Garino,^a Tania Limongi,^a Angelica Chiodoni,^b and Valentina Cauda^a

^a *Dipartimento di Scienza Applicata e Tecnologia, Politecnico di Torino, Corso Duca degli Abruzzi 24, 10129-Torino, Italy*

^b *Center for Sustainable Future Technologies CSFT@Polito, Istituto Italiano di Tecnologia, Corso Trento 21, 10129-Torino, Italy*
E-mail: francesca.susa@polito.it

In the last few years there has been an increasing interest in nanomedicine and in particular in developing nanoparticles to fight cancer [1].

Zinc oxide nanocrystals (ZnO NCs), thanks to their peculiar properties, can be employed for cancer diagnosis and therapy. Even if the cytotoxicity mechanism of ZnO NCs is not totally understood yet, it is most probably due to the combination of intracellular Zn²⁺ cations release and the production of reactive oxygen species: in vitro tests have also shown an increased cytotoxic effect of ZnO on cancer cells with respect to the healthy counterpart [1].

In order to promote the stability of ZnO NCs in physiological environment, increase their biocompatibility and reduce their immunogenic effects, we covered ZnO NCs with a biomimetic phospholipidic bilayer [2] derived from extracellular vesicles, in particular exosomes, obtaining a nanoconstruct called TrojaNanoHorse (TNH).

Exosomes are naturally produced by many types of cells as vehicle of intercellular communication and, when autologous, they are not recognized by the immune system. We extracted exosomes from living cells and we re-used them as a biomimetic, nature-derived coating to stabilize and reduce the immunogenicity of the ZnO NCs. Most importantly, we would like to exploit the natural communication function of exosome to drive the therapeutic nanoparticles towards the cancer cells [3].

We optimized the coupling process between ZnO NCs and exosomes through a systematic study of the thermodynamic, kinetic and electrostatic parameters of the process and then we tested the efficiency of coupling by combining different characterization techniques.

These experiments on the TNH preparation are the starting points to further study the TNH internalization process, mechanisms of causing cell damages, stability in biological fluids, and targeting mechanisms, to make TNH a new theranostic nanoconstruct against cancer.

[1] L. Racca, M. Canta, B. Dumontel, A. Ancona, T. Limongi, N. Garino, M. Laurenti, G. Canavese, V. Cauda, *Smart Nanoparticles for Biomedicine* **1** (2018) 171-187.

[2] B. Dumontel, M. Canta, H. Engelke, A. Chiodoni, L. Racca, A. Ancona, T. Limongi, G. Canavese, V. Cauda, *J. Mater. Chem. B* **5** (2017) 8799-8813.

[3] M.P. Zaborowski, L. Balaj, X.O. Breakefield, C.P. Lai, *Bioscience* **65** (2015) 783-797.